UNITED STATES DISTRICT COURT DISTRICT OF MASSACHUSETTS

JOHN HANCOCK LIFE INSURANCE)	
COMPANY, JOHN HANCOCK	Ś	
VARIABLE LIFE INSURANCE)	
COMPANY, and MANULIFE)	
INSURANCE COMPANY (f/k/a)	Civil Action No. 05-11150-DPW
INVESTORS PARTNER LIFE)	
INSURANCE COMPANY),)	
)	
Plaintiffs,)	
)	
V.)	
)	
ABBOTT LABORATORIES,)	
)	
Defendant.)	

AFFIDAVIT OF OZGE GUZELSU IN SUPPORT OF MOTION FOR PROTECTIVE ORDER REGARDING DEPOSITION TOPICS 1 AND 2 OF HANCOCK'S RULE 30(b)(6) DEPOSITION NOTICE

- I, Ozge Guzelsu, hereby state that:
- 1. I currently am employed as an associate at Munger, Tolles & Olson LLP. I submit this declaration in support of Defendant Abbott Laboratories' ("Abbott's") Motion for Protective Order Regarding Deposition Topics 1 and 2 of Hancock's Rule 30(b)(6) Deposition Notice. If called as a witness, I could and would testify competently to the facts stated herein.
- 2. Attached hereto as Exhibit A are true and correct copies of .excerpts from the deposition of Dr. Bruce McCarthy taken September 29, 2006.

- 3. Attached hereto as Exhibit B are true and correct copies of excerpts of the deposition of Dr. Bruce McCarthy taken March 16, 2007.
- 4. Attached hereto as Exhibit C are true and correct copies of excerpts of the deposition of Dr. Christopher Silber taken February 9, 2007.
- 5. Attached hereto as Exhibit D are true and correct copies of excerpts of the deposition of Marilyn Collicott taken September 27, 2006.
- 6. Attached hereto as Exhibit E are true and correct copies of excerpts of the deposition of James Thomas taken April 13, 2007.
- 7. Attached hereto as Exhibit F are true and correct copies of excerpts of the deposition of Michael Meyer taken January 23, 2007.
- 8. Attached hereto as Exhibit G are true and correct copies of excerpts of the deposition of Andrea Landsberg taken February 16, 2007.
- 9. Attached hereto as Exhibit H are true and correct copies of excerpts of the deposition of Dr. Paul Andrews taken April 4, 2007.
- 10. Attached hereto as Exhibit I are true and correct copies of excerpts of the deposition of Dr. Perry Nisen taken November 22, 2006.
- 11. Attached hereto as Exhibit J are true and correct copies of excerpts of the deposition of Dr. Azmi Nabulsi taken January 24, 2007.
- 12. Attached hereto as Exhibit K are true and correct copies of excerpts of the deposition of Diane D'Amico taken October 26, 2006.
- 13. Attached hereto as Exhibit L are true and correct copies of excerpts of the deposition of Diane D'Amico taken November 28, 2006.
 - 14. Attached hereto as Exhibit M are true and correct copies of excerpts of the

deposition of Jim Looman taken February 1, 2007.

- 15. Attached hereto as Exhibit N are true and correct copies of excerpts of the deposition of Elizabeth Kowaluk taken October 10, 2006.
- 16. Attached hereto as Exhibit O are true and correct copies of excerpts of the deposition of Lise Loberg taken February 2, 2007.
- 17. Attached hereto as Exhibit P are true and correct copies of excerpts of the deposition John Leonard taken November 30, 2006.
- 18. Attached hereto as Exhibit Q are true and correct copies of excerpts of the deposition of Jeffrey Leiden taken April 26, 2007.
- 19. Attached hereto as Exhibit R are true and correct copies of excerpts of the deposition of Philip Deemer taken October 27, 2006.
- 20. Attached hereto as Exhibit S is a true and correct copy of Hancock's Revised Notice of Deposition pursuant to Rule 30(b)(6), dated March 26, 2007.
- 21. Attached hereto as Exhibit T is a true and correct copy of Hancock's Notice of Deposition pursuant to Rule 30(b)(6), dated March 30, 2007.
- 22. Attached hereto as Exhibit U are true and correct copies of excerpts of the deposition of Keith Hendricks taken April 27, 2007.
- 23. Attached hereto as Exhibit V is a true and correct copy of a letter from Jeffrey Weinberger, Esq. to Brian Davis, Esq., dated April 13, 2007.
- 24. Attached hereto as Exhibit W is a true and correct copy of a letter from Joseph Zwicker, Esq. to Jeffrey Weinberger, Esq. dated April 18, 2007.

I declare under penalty of perjury under the laws of the United States of America that the foregoing is true and correct and that this affidavit is executed this 7th day of May, 2007, in Los Angeles, California.



CERTIFICATE OF SERVICE

I hereby certify that this document(s) filed through the ECF system will be sent electronically to the registered participants as identified on the Notice of Electronic Filing (NEF) and paper copies will be sent to those indicated as non registered participants on May 7, 2007.

Date: May 7, 2007.	
	/s/ Michael S. D'Orsi
	Michael S. D'Orsi

Exhibit A, Filed Under Seal

Exhibit B, Filed Under Seal

Exhibit C, Filed Under Seal

Exhibit D, Filed Under Seal

Exhibit E, Filed Under Seal

Exhibit F, Filed Under Seal

Exhibit G, Filed Under Seal

Exhibit H, Filed Under Seal

Exhibit I, Filed Under Seal

Exhibit J, Filed Under Seal

Exhibit K, Filed Under Seal

Exhibit L, Filed Under Seal

Exhibit M, Filed Under Seal

Exhibit N, Filed Under Seal

Exhibit O, Filed Under Seal

Exhibit P, Filed Under Seal

Exhibit Q, Filed Under Seal

Exhibit R, Filed Under Seal

EXHIBIT S

UNITED STATES DISTRICT COURT FOR THE DISTRICT OF MASSACHUSETTS

)
JOHN HANCOCK LIFE INSURANCE	ý
COMPANY, JOHN HANCOCK)
VARIABLE LIFE INSURANCE)
COMPANY, and MANULIFE)
INSURANCE COMPANY (f/k/a)
INVESTORS PARTNER INSURANCE)
COMPANY),) CIVIL ACTION NO. 05-11150-DPW
)
Plaintiffs,)
)
v.)
)
ABBOTT LABORATORIES,)
)
Defendant.)
)

REVISED NOTICE OF DEPOSITION

PLEASE TAKE NOTICE that, pursuant to Fed. R. Civ. P. 30(b)(6), plaintiffs John Hancock Life Insurance Company, John Hancock Variable Life Insurance Company, and Manulife Insurance Company (f/k/a Investors Partner Insurance Company) (collectively, "John Hancock") will take the deposition of defendant Abbott Laboratories on Friday, April 27, 2007, commencing at 9:30 a.m. at the offices of Levenfeld Pearlstein, LLC, 2 North LaSalle Street, Suite 1300, Chicago, Illinois, or such other location as may be mutually agreed to by the parties. Abbott shall designate, prepare and produce one or more knowledgeable officers, directors, or other representatives to testify on its behalf as to the topics set forth below.

PLEASE TAKE FURTHER NOTICE that the deposition noticed above will be recorded stenographically, and through real-time court reporting, such as by LiveNote. The deposition also may be recorded by audio or video technology, such as videotape. The deposition will be taken before a notary public or other person authorized to administer oaths and will continue from day-to-day until completed, Saturdays, Sundays and holiday excepted.

Definitions

For purposes of this Notice, John Hancock adopts the "Uniform Definitions in Discovery Requests" contained in Local Rule 26.5. The following additional terms shall have the meanings set forth below:

- "You," "your" and "Abbott" shall mean defendant Abbott Laboratories, its 1. various corporate parents, subsidiaries, affiliates, subdivisions and departments, and any and all representatives, successors, assigns, officers, directors, employees, agents, attorneys or other persons or entities who have acted or purported to act for or on behalf of any of them.
- "John Hancock" shall mean collectively defendants John Hancock Life Insurance 2. Company, John Hancock Variable Life Insurance Company, and Manulife Insurance Company (f/k/a Investors Partner Life Insurance Company), their various subsidiaries, affiliates, divisions and departments, and any and all representatives, successors, assigns, officers, directors, employees, agents, auditors, attorneys or other persons or entities who have acted or purported to act for or on behalf of any of them, including, without limitation, representatives of the StoneTurn Group.
- The "Research Funding Agreement" shall mean the Research Funding Agreement 3. by and between Abbott and John Hancock, dated as of March 13, 2001.

- 4. The "Program Compounds" shall have the meaning set forth in the Research Funding Agreement.
 - 5. "Regarding" shall have the same meaning as "concerning."
 - 6. "Any" also shall mean "ali," and "all" also shall mean "any."

Topics Of Examination

- 1. Abbott's usual policies, practices, procedures and methodologies, as of 2000 and 2001, for projecting future sales and revenues for the Program Compounds or other pharmaceutical compounds under development by Abbott, including, but not limited to:
 - a. how Abbott considered or analyzed market opportunities for such compounds;
 - b. how Abbott considered or analyzed actual or potential competing products for such compounds;
 - c. any market data or other information considered by Abbott in projecting sales and revenues for such compounds;
 - d. how Abbott considered or analyzed the likelihood of regulatory success for such compounds;
 - e. how Abbott considered or analyzed commercialization costs, such as manufacturing and marketing costs, for such compounds;
 - f. how Abbott considered or analyzed potential profits for such compounds;
 - g. the identity and responsibilities of the persons who had primary responsibility within Abbott for projecting future sales and revenues for such compounds; and
 - h. any other factor that Abbott considered or analyzed in projecting future sales and revenues for such compounds.

- 2. The specific policies, practices, procedures and methodologies utilized by Abbott to project future sales and revenues for the Program Compound known as ABT-518 in 2000 and 2001, including, but not limited to, the policies, practices, procedures and methodologies utilized by Abbott to prepare the sales projections set forth in the documents attached hereto as Exhibit A.
- The specific policies, practices, procedures and methodologies utilized by Abbott to project future sales and revenues for the Program Compound known as ABT-594 in 2000 and 2001, including, but not limited to, the policies, practices, procedures and methodologies utilized by Abbott to prepare the sales projections set forth in the documents attached hereto as Exhibit B.
- 4. The specific policies, practices, procedures and methodologies utilized by Abbott to project future sales and revenues for the Program Compound known as ABT-773 in 2000 and 2001, including, but not limited to, the policies, practices, procedures and methodologies utilized by Abbott to prepare the sales projections set forth in the documents attached hereto as Exhibit C.

JOHN HANCOCK LIFE INSURANCE COMPANY, JOHN HANCOCK VARIABLE LIFE INSURANCE COMPANY, and MANULIFE INSURANCE COMPANY (f/k/a INVESTORS PARTNER INSURANCE COMPANY)

By its attorneys,

Brian A. Davis (BBO No. 546462)

Joseph H. Zwicker (BBO No. 560219) Karen Collari Troake (BBO No. 566922)

Richard C. Abati (BBO No. 651037) Stacy Blasberg (BBO No. 657420)

CHOATE, HALL & STEWART LLP

Two International Place

Boston, Massachusetts 02110 Tele: (617) 248-5000 Fax: (617) 248-4000

Date: March 26, 2007

CERTIFICATE OF SERVICE

I hereby certify that a copy of the foregoing document was served by electronic and overnight mail upon Peter E. Gelhaar, Esq., Donnelly, Conroy & Gelhaar, LLP, One Beacon Street, 33rd Floor, Boston, MA 02108, and Gregory D. Phillips, Esq., Munger, Tolles & Olson LLP, 355 South Grand Avenue, Los Angeles, CA 90071, on this 26th day of March, 2007.

Richard C. Abati

EXHIBIT A

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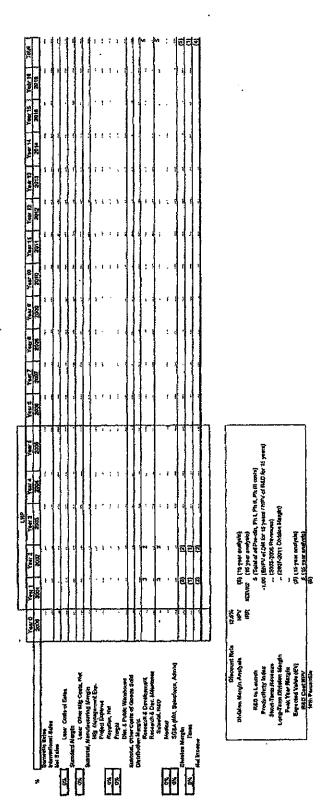
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EXHIBIT T

UNITED STATES DISTRICT COURT FOR THE DISTRICT OF MASSACHUSETTS

JOHN HANCOCK LIFE INSURANCE COMPANY, JOHN HANCOCK VARIABLE LIFE INSURANCE COMPANY, and MANULIFE INSURANCE COMPANY (f/k/a INVESTORS PARTNER INSURANCE)))))
COMPANY),	CIVIL ACTION NO. 05-11150-DPW
- Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Contain the Cont)
Plaintiffs,)
)
v.)
)
ABBOTT LABORATORIES,)
)
Defendant.)

NOTICE OF DEPOSITION

PLEASE TAKE NOTICE that, pursuant to Fed. R. Civ. P. 30(b)(6), plaintiffs John Hancock Life Insurance Company, John Hancock Variable Life Insurance Company, and Manulife Insurance Company (f/k/a Investors Partner Insurance Company) (collectively, "John Hancock") will take the deposition of defendant Abbott Laboratories on April 12, 2007 commencing at 9:30 a.m. at the offices of Levenfeld Pearlstein, LLC, 2 North LaSalle Street, Suite 1300, Chicago, Illinois, or such other location as may be mutually agreed to by the parties. Abbott shall designate, prepare and produce one or more knowledgeable officers, directors, or other representatives to testify on its behalf as to the topics set forth below.

PLEASE TAKE FURTHER NOTICE that the deposition noticed above will be recorded stenographically, and through real-time court reporting, such as by LiveNote. The deposition also may be recorded by audio or video technology, such as videotape. The deposition will be taken before a notary public or other person authorized to administer oaths and will continue from day-to-day until completed, Saturdays, Sundays and holidays excepted.

Definitions

For purposes of this Notice, John Hancock adopts the "Uniform Definitions in Discovery Requests" contained in Local Rule 26.5. The following additional terms shall have the meanings set forth below:

- 1. "You," "your" and "Abbott" shall mean defendant Abbott Laboratories, its various corporate parents, subsidiaries, affiliates, subdivisions and departments, and any and all representatives, successors, assigns, officers, directors, employees, agents, attorneys or other persons or entities who have acted or purported to act for or on behalf of any of them.
- 2. "Abbott's Senior Management" shall mean the Abbott personnel who had or have overall responsibility, authority and accountability for managing Abbott's Global Pharmaceutical Research and Development organization and operations, including, without limitation, Miles D. White and the "Senior Management" referenced in Abbott Document No. ABBT0101924.
- 3. "John Hancock" shall mean collectively defendants John Hancock Life Insurance Company, John Hancock Variable Life Insurance Company, and Manulife Insurance Company (f/k/a Investors Partner Life Insurance Company), their various subsidiaries, affiliates, divisions and departments, and any and all representatives, successors, assigns,

officers, directors, employees, agents, auditors, attorneys or other persons or entities who have acted or purported to act for or on behalf of any of them, including, without limitation, representatives of the StoneTurn Group.

- The "Research Funding Agreement" shall mean the Research Funding 4. Agreement by and between Abbott and John Hancock, dated as of March 13, 2001.
- 5. The "Program Compounds" shall have the meaning set forth in the Research Funding Agreement.
- 6. "Program Term" shall have the meaning set forth in the Research Funding Agreement.
 - "Regarding" shall have the same meaning as "concerning." 7.
 - "Any" also shall mean "all," and "all" also shall mean "any." 8.
- "And" as well as "or" shall be construed both disjunctively and conjunctively to 9. mean "and/or."

Topics Of Examination

- Abbott's knowledge and belief concerning the prospects and condition (including 1. safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-518 as of March 13, 2001.
- 2. Abbott's knowledge and belief concerning the prospects and condition (including safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-594 as of March 13, 2001.

- Abbott's knowledge and belief concerning the prospects and condition (including 3. safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-773 as of March 13, 2001.
- The knowledge and belief of each member of Abbott's Senior Management 4. concerning the prospects and condition (including safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-518 as of March 13, 2001.
- 5. The knowledge and belief of each member of Abbott's Senior Management concerning the prospects and condition (including safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-594 as of March 13, 2001.
- The knowledge and belief of each member of Abbott's Senior Management 6. concerning the prospects and condition (including safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-773 as of March 13, 2001.
- 7. Abbott's valuation of, and methods for valuing (including, without limitation, any models used in such valuations) the Program Compound known as ABT-518 at any time from January 1, 2001 to the present.
- 8. Abbott's valuation of, and methods for valuing (including, without limitation, any models used in such valuations) the Program Compound known as ABT-594 at any time from January 1, 2001 to the present.
- 9. Abbott's valuation of, and methods for valuing (including, without limitation, any models used in such valuations) the Program Compound known as ABT-773 at any time from January 1, 2001 to the present.
- Abbott's nominal or intended and reasonably expected spending on the Program 10. Compound known as ABT-518 at any time from January 1, 2001 to the present.

- 11. Abbott's nominal or intended and reasonably expected spending on the Program Compound known as ABT-594 at any time from January 1, 2001 to the present.
- 12. Abbott's nominal or intended and reasonably expected spending on the Program Compound known as ABT-773 at any time from January 1, 2001 to the present.
- 13. Abbott's reasons for discontinuing or terminating the development of the Program Compound known as ABT-518.
- 14. Abbott's reasons for discontinuing or terminating the development of the Program Compound known as ABT-594.
- 15. Abbott's reasons for discontinuing or terminating the development of the Program Compound known as ABT-773.
- 16. All communications among or between Abbott's Senior Management, at any time from January 1, 2000 to the present, regarding the prospects or condition (including safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-518.
- 17. All communications among or between Abbott's Senior Management, at any time from January 1, 2000 to the present, regarding the prospects or condition (including safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-594.
- 18. All communications among or between Abbott's Senior Management, at any time from January 1, 2000 to the present, regarding the prospects or condition (including safety, efficacy, scientific viability or commercial viability) of the Program Compound known as ABT-773.

- 19. All communications among and between senior Abbott's Senior Management, at any time from January 1, 2000 to the present, regarding the actual or potential discontinuation or termination of development of the Program Compound known as ABT-518.
- 20. All communications among and between senior Abbott's Senior Management, at any time from January 1, 2000 to the present, regarding the actual or potential discontinuation or termination of development of the Program Compound known as ABT-594.
- 21. All communications among and between senior Abbott's Senior Management, at any time from January 1, 2000 to the present, regarding the actual or potential discontinuation or termination of development of the Program Compound known as ABT-773.
- 22. The creation of, analysis reflected in, or actions taken by Abbott in connection with, the document attached hereto as Exhibit A as it pertains to the Program Compounds known as ABT-518, ABT-594 and ABT-773.
- 23. The creation of, analysis reflected in, or actions taken by Abbott in connection with, the document attached hereto as <u>Exhibit B</u> as it pertains to the Program Compounds known as ABT-518, ABT-594 and ABT-773.
- 24. The creation of, analysis reflected in, or actions taken by Abbott in connection with, the document attached hereto as Exhibit C as it pertains to the Program Compound known as ABT-594.

JOHN HANCOCK LIFE INSURANCE COMPANY, JOHN HANCOCK VARIABLE COMPANY, and LIFE INSURANCE INSURANCE COMPANY MANULIFE (f/k/a INVESTORS PARTNER INSURANCE COMPANY)

By its attorneys,

Brian A. Davis (BBO No. 546462)

Joseph H. Zwicker (BBO No. 560219)

Karen Collari Troake (BBO No. 566922)

Richard C. Abati (BBO No. 651037)

CHOATE, HALL & STEWART LLP

Two International Place

Boston, Massachusetts 02110

Tele: (617) 248-5000 Fax: (617) 248-4000

Date: March 30, 2007

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CERTIFICATE OF SERVICE

I hereby certify that a copy of the foregoing document was served by electronic and overnight mail upon Peter E. Gelhaar, Esq., Donnelly, Conroy & Gelhaar, LLP, One Beacon Street, 33rd Floor, Boston, MA 02108, and Gregory D. Phillips, Esq., Munger, Tolles & Olson LLP, 355 South Grand Avenue, Los Angeles, CA 90071, on this 30th day of March, 2007.

Exhibit A

TIAL PORT	FOLIO	PRIORITIZATION		C- continue P- pending T- terminate
Project	Priority	Next steps	Responsibility	Tinding
Anti-Infectives ABT-492	¢	Address safety issues (including QTc) with internal/ expert review Determine how many indications at launch (pey back)	+ J. Leonard	-
H\$R-903	٣	Consider trading with Dalichi Halt any new expenditure	• J. Tyree	-
ABT-773	С	Assess side effects issues with expert review (QTc and liver tox.) Ensure all drug interactions are adequately covered Assess relative to Ketek	J. Leonard J. Leonard I. Leow	•
Urolegy BSF 420627	e q .	Sot up task force to address issues and bring back plan to senior management Heasons for failure of the SKB ETa/b aniagonist Design short (-4 week) PoP trial for symptom relief Hallonale for sustained release formulation Nature of the Schwarz Phoma relationable	• J. Leonard	+ By May
Hypothyroidism T3/T4	P	Assess most appropriate ratio Gain FDA feedback on study design Determine ex-US market attractiveness (price)	• J. Leonard	• Ву Мау
Asthma Hokunalin lape	P	Conduct market research on acceptance by different patient segments Determine how to position against long acting beta against and combination inhalers Evaluate apportunity to gain complete access to the	• A. Higgins/ E. Florensino • J. Tyree	• May

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TIAL POF	RTFOLIO	PRIORITIZATION (CONTINUED)		C- continu P- pending T- termina
Project	Priority	Next steps	Responsibility	Timing
Oncology ABT-510	С	Pursus proof of concept Leverage TAP knowledge of anglogenesis product development (appropriate endpoints)	• Project team	- As planned
ABT-761	¢	Pursue proof of concept Use scinocadiogram to monitor potential cardiotoxidity Resolve potent drug menujacturing approach	Project leam CNC group	• As planned
ABT-518	Hold	Well for May results from Plizer (will save ~\$1 mill) and re-evaluate Helt all luther expenditure	Serior management	• May
Rubliecan	Ρ .	Significant clinical rework required (funded by partner)- further in-depth review required Make a proceed decision when 2Q data available	• J. Leonard	• By May
Theragyn	Р	Negative initial scientific perspective - further in- depth review required, e.g., Cetamithe if there is a PoC to support claim Address GMP issues Determine best control to demonstrate officecy Re-look at partnership contract	 J. Leonald J. Tysee 	By May By May
ABT-627	c	Seek alternative funding (e.g., NOI) before starting major trial If move shead Determine how to ensure NDA filing in 2004 Get FDA input since survival not primary endpoint Harmonize US and EU study design and inputs Consider partnership (e.g., Bi or established oncology player)	• J. Leonard, P. Nisen • J. Tyree	• ASAP

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Project	Priority	Next steps	Responsibility	Timing
Cardiology/ thrombosis Darbsenian (LU 135252)	Hold	Continue currently budgeted funding for next 5 months Do not start any new trials (e.g., hypertension planned for May)	• Project team	+ Ongoing
		If proceed, plan for pilot to look at effects in sperm and telratogenecity Consider out-license or swap	• J. Tyree	• ASAP
LU 208075	Hold	 Continue currently budgeted funding for next six 	 Project leam 	∗ ongoing
		months + Look at Myogen deal + Out-license or swap	J. Tyree	
Levosimendan	C	 Conduct detailed expert panel review for trial design. 	 J. Leonard 	• May
PEG-hindin	P	 Set up expert panel for commercial assessment (Is diabetes an option?) 	• E. Ogunro	• By May
Ancrod	т	 Identity out-licensing opportunities 	- J. Tyree	• TBD
Urokinase	P	Market research required on open cath Match versus IPA in dose-ranging studies to determine efficacy	+ E. Florenilno	• By May
Pro-urokinase	Ç	 Identify apportunities to speed up program 	 Project team 	• TBD
Clivarine	C	 Assessment by HPD (review previous evaluation and new trial data) 	• E. Ogunro	• By May
		 Understand linished product manufacturing cost 	 B. Dempsey 	
Aythmol SR	С	Continue filing Verify II package is likely approvable Assess commercial attractiveness in a generic market	Project leam	 Ongoing

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INITIAL PORTFOLIO PRIORITIZATION (CONTINUED)

C- continue P-pending T-terminate

Project	Priority	Next steps	Responsibility	Timing
Neuroscience ABT 594	P	Await results from ongoing Pli trial — probable T Project team to develop decision criteria for goino go	• Senior management	• June/
ABT 963	С	 Identity a co-development/co-promotion partner (TAP high on list) Evaluate benefits of the long half life in pain and cancer (including additional physician market research) Exprore cancer prophyticats and Alzhelmer's Indications 	• J. Tyres • Project team	• TBD
BSF 201640	P	Complete review of all schizophrenia NCEs with expert panel Complete staffing of internal project learn, but halt further expenditure beyond locking at hepatic lox, and QTC Understand Novarile contract and level of interest	I. Loew Project team J. Tyree	• By May
BSF 190555	P	Complete review as above Halt turther expenditive pending outcome	• I. Loew	• As above
8SF 74398	C	Allow DayCo to continue development Re-look at relationship with DayCo	 Froject team J. Tyree 	• By May
Olluadid Otos	Hold	Return to ALZA or out-license to other interested partner.	۰ J. Tyree	• TBD
Hydrocodone	C	Assess regulatory pathway Understand DEA impact on manufacturing	• Project team	• By May
Bimaciomol (AST 822)	P	Await data from ongoing klai in April before deciding whether to continue - probable T Hatt further expenditure pending outcome	• Senior management	• April

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TULK	PRIORITIZATION (CONTINUED)	C- continue P- pending T- terminate	
Priority	Next steps	Responsibility	Timing
P	Conduct U.S. commercial assessment with TAP Assess how to position in Europe versus generics and	• E. Florensino	• By June
	Develop model to assess spend at different termination points	• Bob Funck	• By May
۲	• Terminate outside Japan	 Project team 	• Immediat
Ċ	Develop and pursue a small PoC trial in humans ASAP (consider niche indication first and leverage Martene's according)	• Project team	- ASAP
	Conduct market research on IBS versus constipation (including pricing)	• E. Ptorentino	
o	Conduct intensive product review 2 day meeting with J. Lennard's group (already in process)	• J. Leonard	• By May
	Important actions include - Approach FDA for last frack and compassionate use - Develop strategy for DMARD claim in first submission - Assess need for Enbrel assay to detect HAHAs - Assess delivery device options - Evaluate additional indications (e.g., Psortasis, Crohne, heart failure) and pediatric program - Profile Cellitech product - Assess Impact of additional IV program on relimbusement	• Various	• By May
	P T C	P • Conduct U.S. commercial assessment with TAP • Assess how to position in Europe versus garantes and implications for comparative intal • Develop model to assess spend at different termination points T • Terminate outside Japan C • Develop and pursue a small PoC trial in humans ASAP (consider riche indication first and leverage Martene's expertise) • Conduct market research on IBS versus constitution (including pricing) C • Conduct intensive product review • 2 day meeting with J. Lennard's group (already in process) • ½ day session with senfor management group • important actions include • Approach FDA for fast frack and compassionate use • Develop strategy for DMARD claim in first submission • Assess need for Entirel assay to detect HAHAs • Assess delivery device options • Evaluate additional indications (e.g., Psoriasis, Crohne, heart failure) and pediatric program • Profile Cellisch product • Assess Impact of additional IV program on	Priority Next steps

INITIAL PORTFOLIO PRIORITIZATION (CONTINUED)

C- continue P- pending T- terminate

Project	Priority	Next steps	Responsibility	Timing
immunology (continued) Segard	Hold	Continue filing in EU and Canada Put on hold in US – consider creating a small learn in the US to analyse data, propose smaller Pil study Research pricing, marketing and Phase IV plans in Europe Look at TNF-elpha levels retrospectively to see stratification with IL-B Assess manufacturing strategy Identity potential out-licensing opportunities (Generatech)	• Project team • J. Leonard • J. Tyree	• Ongoing
J8 9 5	p	Deckle on most attractive indications from Abboit and pariner perspective Discuss with partner ways to share the various indications and potential for TNF-alphe combinations Add commercial person to the project team by this week	• E, Florentino • J. Tyree • Ongoing	• ASAP

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INITIAL PORTFOLIO PRIORITIZATION (CONTINUED)

C-continue P-pending T-terminate

Project	Priority	Next ateps	Responsibility	भूगांकार्
PIV programs Clarithromycln	c	+ None idenlified	•	-
Omnicel	G	Talk to partners	• J. Tyree	•
Kaletra	С	None identified	• •	-
Norvir	C	None Identified	-	•
Meridia	Hold	Conduct commercial assessment for CNS and depression (P&L)	• B. Dempsey, J. Amott, E. Floreplino	* ASAP .
		 Assess combination therapy with librales Assess outcomes that design to meet preferred commercial profile; determine payback 	• Project team	
Uprima	С	 Ensure no redundant trials with TAP in Europa 	 Project team 	 Ongoing
Trandolapni paich	T	• Halt all activities	- Project team	 immediate
Tranciolapril "Invest" clinical program	P	 Review that in more detail (reduce complexity and risk) 	+ E. Florentio	• By May
Other trandolapril Mais	C	 Continue "Create", "Peace" and "Benedict" Indi programs 	• Project team	 Ongoing
Fenolibrate	С	 Develop co-formulation ideas with Meridia and staths (including assessment of sales and costs) 	• Project learn	•
Depakote	C	None Identified	•	•
Gengral	C	None identified	-	-

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Exhibit B

TIAL PORT	FOLIO	PRIORITIZATION		C- continue P- panding T- terminal
Project	Priority	Next stops	Responsibility	Timing
Anti-infectives ABT-492	С	Address safety issues (including QTo) with internal/ expert review Determine how many indications at leunch (pay back)	• J. Leonard	•
HSR-903	T	Consider trading with Dalichi Halt any new expenditure	• J. Tyree	•
ABT-773	C	Assess side effects issues with expert review (QTc and liver tox.) Ensure all drug interactions are adequately covered Assess retailve to Ketek	J. Leonard J. Leonard I. Leow	•
Urology BSF 420627	Р	Set up tack force to address issues and bring back plan to senior management Reasons for failure of the SKB ETa/b antagorist Design short (~4 week) PoP trial for symptom relief Rationale for sustained release formulation Nature of the Schwarz Pharma relationship	• J, Leonard	+ By May
Hypothyroldism T3/T4	P	Assess most appropriate ratio Gain FDA feedback on sludy design Determine ex-US market attractiveness (price)	⊮J. Leonard	• Sy May
Asthma Hokunalin tape	P	Conduct market research on soceptance by different patient segments Determine how to position against long acting beta agonists and combination inhalers Evaluate opportunity to gain complete access to the patch technology	A. Higgins/ E. Florentino J. Tyree	• May

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TIAL POF	RTFOLIO	PRIORITIZATION (CONTINUED)		C- continue P- pending T- terminal
Project	Priority	Next steps	Responsibility	Timing
Oncolegy ABT-510	c	Pursue proof of concept Leverage TAP knowledge of anglegenests product development (appropriate endpoints)	• Project learn	• As planned
ABT-751	C	Pursue proof of concept Use echocardiogram to monitor potential cardiotoxicity Resolve potent drug manufacturing approach	Project learn CMC group	 As planned
ABT-518	Hold/T	Well for May results from Pfizer (will save ~\$1mill) and re-evaluate Hait all further expenditure	Sentor management	- May
Rubliecan	5	 Significant clinical rework required (funded by partner)- furthor in-depth review required Make a proceed decision when 2Q data available 	• J. Leonard	• By May
Theragyn	P	Negative Initial scientific perspective - kuther Indepth review required, e.g., Determine if there is a PoC to support claim Address GMP issues Determine best control to demonstrate efficacy Re-look at partnership confired.	 J. Leonard J. Tyree 	• By May • By May
ABT-627	o	Seek allamative funding (e.g., NCI) before starting major trial If move ahead Determine how to ensure NDA filing in 2004 Get FDA Input since survival not primary endpoint Harmonize US and EU study design and imputs Consider partnership (e.g., B) or established oncology player)	J. Leonard, P. Nisen J. Tyree	• By May

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TIAL POR	TFOLI	O PRIORITIZATION (CONTINUED)		C- continue P- pending T- terminal
Project	Priority	Next steps	Responsibility	Timing
Cardiology/ thrombosis				
Darusenian (LIJ 135252)	Hold/Y	Continue currently budgeted furning for next 6 months Do not start any new trials (e.g., hypertension planned The start	 Project team 	Ongoing
		lor May) Consider out-license or swap	• J. Tyree	+ ASAP
LU 208075	Hotd/T	Continue currently budgeted funding for next six months	• Project team	 ongolng
		Look at Myogen deal Out-license or swep	J. Tyree	
Levosimendan	C	- Conduct detailed expert panel review for trial design	• J. Leonard	• May
PEG-hirudin	Р	 Set up expert panel for commercial assessment (le diabetes an option?) 	E, Ogunio	• By May
Ancrod	T	 identity out-licensing opportunities 	J. Tyree	TBD
Urokinase	P	Market research required on open callh Match versus IPA in dose-ranging studies to determine stillosoy	• E. Florentino	• Ву Мау
Pro-urckinase	¢	* Identity opportunities to speed up program	 Project team 	• TBD
Clivatine	C	Assessment by HPD (review previous evaluation and new trial data)	・ 性. Ogunro	• By May
		 Understand finished product manufacturing cost 	 B. Dempsey 	
Rythmol SR	C	Continue filing Verify if package is likely approvable Assess commercial altractiveness in a generic market	• Project leam	Ongoing

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INITIAL PORTFOLIO PRIORITIZATION (CONTINUED)

C- continue P-pending T-terminate

Project	Priority	Next steps	Responsibility	Timing
Neuroscience ABT 594	Р	Awali results from ongoing Pil trial probable T Project learn to develop decision criteria for go/no go	• Sanior management	+ June/ July
ABT 963	C	identify a co-development/co-promotion partner (TAP high on list) Evaluate banefils of the long half lite in pain and cancer (including additional physician market research) Explore cancer prophylaxis and Alzhelmer's indications	J. Tyres Project leam	• TBD
BSF 201640	Þ	Complete review of all schizophrenia NCEs with expert panel Complete statting of internal project team, but helt further expenditure beyond looking at hapatic tox, and QTc Understand Novarits contract and level of interest	I, LoewProject leamJ. Tyree	+ By May
BSF 190555	P	Complete review as above Hait further expenditure pending outcome	• f. Loew	• As above
BSF 74398	C (no cost)	Allow DevCo to confirme development Re-look at relationship with DevCo	 Project team J. Tyrea 	• By May
Dilundid Oros	Hold/T	Return to ALZA or out-license to other interested partner	J. Tyres	• TBD
Hydrocodone	C	Assess regulatory pathway Understand DEA Impact on manufacturing	• Project learn	• By May
Bimaciomol (ABT 822)	۶	Await data from ongoing trial in April before deciding whether to confinue - probable T Halt further expenditure pending autoome	 Senior management 	• Aprili

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ITIAL PORT	FOLIC	PRIORITIZATION (CONTINUED)	C- continue P- pending T- terminate	
Project	Priority	Next steps	Responsibility	Timing
Gastro-enterology Ganaton	p	Conduct U.S. commercial assessment with TAP Assess how to position in Europe versus generics and implications for comparative trial	- E. Florentino	• By June
		Develop model to assess spend at different termination points	- Bob Punck	• By May
TU-198	Τ	Terminate outside Japan	• Project learn	• immediate
AU-224	С	 Develop and pursue a small PoC trial in humans ASAP (consider riche indication first and leverage Mariete's expedise) 	• Project leam	- ASAP
		 Conduct market research on IBS versus constipation (including pricing) 	• E. Florentino	
immunology				B
D2E7	¢	 Conduct intensive product review 2 day meeting with J. Lennard's group (already in process) 16 day session with senior menagement group 	+J, Leonard	• By May
		- Typoriant actions include - Approach FDA for last track and compassionate use - Develop strategy for DMARD claim in first submission - Assess need for Erdytel assay to detect HAHAS - Assess delivery device options - Evaluate additional indications (e.g., Psoriasis, Crohns, heart failure) and pediatric program - Profile Cellisch product - Assess impact of additional IV program on reimburgement	• Various	• By May
		 Develop list of potential marketing partners for quids 	• J. Tyree	

INITIAL PORTFOLIO PRIORITIZATION (CONTINUED)

C- continue P- pending T- terminate

Project	Priority	Next steps	Responsibility	Timing
immunology (continued) Segard	Hold	Continue Illing in EU and Canada Put on hold in US – consider creating a small team in the US to analyse data, propose smaller Pil study Research pricing, marketing and Phase IV plans in Europe Look at TNF-alpha levets retrospectively to see stratication with IL-6 Assess manufacturing strategy	• Project team • J. Leonard	Ongoing
		- locality potential out-licensing opportunities (Generalech)	• J. Tyree	
J695	P	 Decide on most attractive indications from Abbott and partner perspective 	• E. Florentino	• ASAP
		Discuss with partner ways to share the various indications and potential for TNF-alpha combinations	• J. Tyree	
		· Add commercial person to the project learn by this week	 Ongoing 	

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INITIAL PORTFOLIO PRIORITIZATION (CONTINUED)

C- continue P- pending T- terminate

Project	Priority	Next steps	Responsibility	Timing
PiV programs Clarithromych	0	Nane Identified	-	-
Omnicef	C	None identified	-	•
Kaletra	C	 None identified 	•	-
Norvir	Ç	None identified	•	•
Meridia	Hold	Conduct commercial assessment for CNS and depression (P&L)	 B. Dempsey, J. Amolt, E. Fiorentino 	• ASAP
		 Assess combination therapy with fibrates Assess outcomes trial design to meet preferred commercial profile; determine payback 	• Projeci team	
Uprima	C	· Ensure no redundant trials with TAP in Europe	• Project leam	 Ongoing
Trandolaprii patch	۲	* Halt all activities	→ Project lears	+ Immediate
Trandolapril "Invest" clinical program	P	 Review trial in more detail (reduce complexity and risk) 	• E. Fiorenlio	- By May
Other trandotapril Irials	C	 Continue "Create", "Pesce" and "Banedict" trial programs 	 Project team 	• Ongoing
Fenolityale	¢	 Develop co-formulation ideas with Meridia and statiss (including assessment of sales and costs) 	• Project team	•
Depakote	C	None Identified	•	•
Gengrali	Ç	None (den)illed		*

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Exhibit C

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NO. 1275 P. 20

Barbara Massa

Mar Icen

Mike Comilla

Matt Russell Tom Woldat

Steve Cohen Mike Higgins

ANALGESIA VENTURE

2001 PLAN Revised 1/26/01 John Leonard Chris Silber George Carter

Bruce McCarthy Mike Biamesen

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-NOV. 20. 2003 8:23AM

NO. 1275 P. 21

Analgesia Venture 2001 PLAN Review (Pass II) Table of Contents

Venture Functional Expense NPS 1776 Project Expense ABT-963 Project Expense ABS-103 Project Expense ABT-089 Project Expense ABT-594 Project Expense NPS 1776 Key Statistics ABS-103 Key Statistics ABT-953 Key Statistics ABT-089 Key Statistics ABT-594 Key Statistics Summary of Projects Blue Plan Summary NPS 1776 Grants ABT-963 Grants 4BS-103 Grants ABT-089 Grants ABT-594 Grants

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Summary	2001 PLAN Pass II
	Summary

	2001 Target	2000 AGU	2001 PLAN	Target vs PLA? Fav(Unfav) Va
ABT - 594	8,900	14,411	6,307	(407)
ABT - 089	}	3,000	613	(619)
NPS 1776	ŧ	:	537	(537)
ABS - 103	:	:	•	ī
ABT - 963		4,000	981'1	(1,186)
Venture Total	8,900	21,411	11,643	(2,743)

a includes a \$120,000 charge from SPD not in Oracle b Completion of work started in 2000, bringing it to a logical holding position. c includes a \$490,000 charge from SPD included in Oracle in error. NOV. 20. 2003 8:24AM

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Analgals Venture
ABT-594
2001 PLAN KEY STATISTICS Pass II
(S000)

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PHARMACEUTICAL PRODUCTS RESEARCH AND DEVELOPMENT 2000 AUGUST UPDATE / 2001 PLAN G0-143010 CCM ABT594 (BASE & ORAL PAIN)

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26-Jan-01	2000	2000	AUG. UPD VS.	2001	PLAN VS.
•	APU	AGU	APR. UPD.	PLAN	AUG, UPD.
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PPD INVESTIGATIONAL DR			/ 2 1\	8.5_	(32)
PPD Investigational Drug QA	23 23	55 55	(32)	86	(32)
	23	22	زهد	•••	4 7
Venture Management		4 400	446	3,988	.505
Analgesia/CCM Venture	4,739	4,493	246 246	3,988	505
	4,739	4,493	240	5,500	
Discovery		*-	mc)	26	24
Advanced Technology	2.5	50	(25)		(51)
Neurological & Urological Res	177	<u></u> 50	(25)	77	(28)
	25	> 0	(44)	•	(,,,,,)
Drug Safety	<i>t</i>		44	187	(118)
Experimental Science	23	70	(46)	409	(120)
Clinical Drug Analysis	290	290	471	233	663
Toxicology	1,366	896 572	471 32	493	79
Pathology	604	591	J.L	34	5\$7
Comparative Medicips	591 4		. 4	7	(7)
Strategic & Exploratory Science	2,877	2,417	460	1,362	1,055
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Pharm Analytical R&D	001	879	(88)	641	238
ANALYTICS DEV & SUPPORT	791 764	745	19	226	519
FORMULATION DEV & SUPPORT	704 403	607	(204)	145	462
CLINICAL FINISHING	197	178	20	63	115
PROJECT MGMT SUPPORT	2,155	2,409		1,075	1,334
PHASE-I CENTER		-,		•	
Phase-I Admin/Pharmacokinetics	185	185		259	(74)
ACPRU	23	25	(2)	367	(343)
ACPRO	208	210	(2)	627	(417)
Development Operations			٠,,		
-	475	475	. 110	259	216
Data Management Statistics	360	171		129	42 .
ABBOTT RES & LIBRARY INF-ARL	89	89		140	(51)
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Regulatory Affairs	,		• •		·
Regulatory Affairs	20	20	***	151	(131)
Research QA	131	80		82	<u></u>
toward Au	151	100		232	(132)
Medical Affairs		-**	•		
Genetics/Admin				2	(2)
Medical Services	53	53		10	43
Outcomes Res/Admin.	42	42		37	5_
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Administration					
R&D Operations/Project Services	75	43	32_	45_	(2)
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25-Jan-01	2000			PLAN	AUG. UPD.	
	apu	AGU	APR. UPD.	- FLAN	·	
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PPD R&D SERVICES PURC	H				-35	
SPD Services Purchased	235	235	2-1		235 235	
ShD 28tAlces Edictingen	235	235	***	14	235	
CLINICAL GRANTS					1 500	
CLINICAL GRANTS	3,000	2,800	200	1,065	1,735	-
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	14,357	13,661	696	9,187	4,474	
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 PROJECT GLOBAL PPD REPORT BY PROJ SUBDIV

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Exhibit U, Filed Under Seal

 $\mathbf{EXHIBIT}\;\mathbf{V}$

MUNGER, TOLLES & OLSON LLP

CARTH T. VINCENT
TED DANE
MARK SHINDERMAN
STUART N. SENATOR
MARTIN D. SERN
DANIEL P. COLUMS
STEVEN S. WEISBURD
EDWARD C. MAGENOTI, JR
RICHARD E. DRIDOYAN
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April 13, 2007

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VIA E-MAIL AND US MAIL

MOLYN HOECKER LUEDTKE Andern Hoecki Mark H. Rum Milison E. Stein

Brian A. Davis Choate, Hall & Stewart LLP Two International Place Boston, Massachusetts 02110

Re: Hancock v. Abbott Laboratories, No. 05-11150-DPW

Dear Brian

This is in response to your letter of April 10, 2007 regarding the parties' respective Rule 30(b)(6) deposition notices.

Hancock's Rule 30(b)6 Notice

On topics 1 and 2, Hancock has already deposed, or is scheduled to depose, the individuals most knowledgeable regarding these topics. As you know, the key individuals who were in charge of both of these programs are former employees. As you recognized in your letter, we do not have the obligation to "seek out and collect information from former personnel" in preparing a Rule 30(b)(6) witness. Any witness that Abbott would now produce on these topics would be required to read this deposition testimony and, to the extent he/she could remember it, repeat it to you. This is a colossal waste of time and resources, and we do not propose to engage in it. However, as I indicated in my April 9 letter, we are willing to designate deposition testimony on these subjects as to which Abbott would agree to be bound in the same manner as if a 30(b)(6) witness was produced.

MUNGER, TOLLES & OLSON LLP Brian A. Davis Two International April 13, 2007 Page 2

> Your letter refers to inconsistencies or other problems with the testimony to date regarding these subjects. If you could be more specific, and identify any particular information that you believe you do not have and which could be provided by a Rule 30(b)6 witness, we will reevaluate our position.

> On topic no. 3, Abbott intends to designate Stanley Bukofzer. Since Dr. Bukofzer is already scheduled to be deposed on April 25 on the same subjects, we expect that you can combine his individual deposition with the deposition under Rule 30(b)(6).

> Abbott designates Dr. John Leonard as its Rule 30(b)(6) representative concerning topics 22 through 23. As you know, Dr. Leonard's deposition has been postponed to a date in May at your request in return for your agreeing to postpone the Blewitt and Klotz depositions. We expect to give you dates for Dr. Leonard after you provide us with dates for Messrs. Blewitt and Klotz. We are looking into the issue of the appropriate witness for topic 24.

Abbott's 30(b)6 Notice

Please provide dates for the Rule 30(b)(6) witnesses you agreed to produce as well as the identity of those witnesses. With respect to topics 5 through 8 of Abbott's notice, we do not need a separate Rule 30(b)(6) deposition for Mr. Friedman because we will be taking his deposition in May anyway. However, please do not interpret this statement as acquiescing to the admission of any expert opinions which were not timely disclosed in accordance with the Federal Rules, as we will vigorously oppose any attempt to do so. Simply stated, regardless of whom you designate, it is our position that neither Mr. Friedman nor any other expert may offer testimony at trial on these subjects because it was not disclosed in expert reports. Of course any "fact" testimony that Mr. Friedman would try to offer would clearly be hearsay.

In connection with requests 10 through 21, to the extent they pertain to Hancock's Committee of Finance, we are entitled to a clear statement from you that Hancock has no present information about the bases or criteria for the Committee of Finance's approval of investments in 2001 or, in the alternative, to a witness who can tell us what this knowledge is.

Finally, with respect to Abbott's topics 22 through 23, we will reserve our rights and wait until your Rule 30(b)(6) witnesses have testified to determine whether to pursue this further. I will note, however, that you are refusing to produce these witnesses on the same basis as stated in our objections to your requests 13 through 21.

einberger

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MUNGER, TOLLES & OLSON LLP Brian A. Davis Two International April 13, 2007 Page 3

bcc:

Peter N. Witty Greg S. Phillips Eric J. Lorenzini Ozge Guzelsu Kathryn Herrity Sonia Arteaga

EXHIBIT W

April 18, 2007

Joseph H. Zwicker (617) 248-5065 jzwicker@choate.com

BY ELECTRONIC AND OVERNIGHT MAIL

Jeffrey I. Weinberger, Esquire MUNGER, TOLLES & OLSON LLP 355 South Grand Avenue, 35th Floor Los Angeles, CA 90071

Re:

John Hancock Life Insurance Company, et al.

v. Abbott Laboratories

U.S.D.C. (Mass.) Civil Action No. 05-11150-DPW

Dear Jeff:

I am responding to your letter of April 13, 2007 to Brian Davis which we received after the close of business on Friday. Brian is out of the office this week. I understand that we are attempting to schedule a call later today that may resolve some outstanding scheduling issues.

Hancock's Rule 30(b)(b) Notice

Jeff, we are entitled to Abbott's knowledge regarding topics 1 and 2. Thus far, the testimony from the individual deponents on these subjects may fairly be characterized as incomplete. Persons from whom we would have expected to obtain relevant testimony, such as Bruce McCarthy, Chris Silber, and Phil Deemer, regularly claimed a lack of recollection of key events that were codified in documents they authored. Perhaps John Leonard is the appropriate choice for these topics. I'll leave that decision to Abbott.

We're amenable to combining Bukofzer's individual and 30(b)(6) testimony assuming we're not going to be limited to the 7 hours for both events. Unfortunately, for several reasons (including that I've just been notified that I have to argue a preliminary injuction motion in Springfield, MA on April 25), the April 25 date needs to be moved back a few days. We can discuss that later.

We're working on dates for Blewitt, Klotz, and Friedman and will attempt to schedule them as you requested for May 8-10.

Letter to Jeffrey I. Weinberger, Esq. MUNGER, TOLLES & OLSON LLP April 18, 2007 Page 2

Abbott's 30(b)(6) Notice

For topics 10-21, we plan to designate Willma Davis regarding issues related to the Bond Investment Committee. We propose Ms. Davis' deposition here in Boston on the morning of April 24, 2007. Ms. Davis has some half days available that week but I would not expect that her deposition would take more than a morning. I am continuing to work with Hancock to discover whether there is information relating to the activities of the Finance Committee as set forth in your Notice.

Other Depositions

Brian will shortly propose dates for Leonard and Tucker. Can you give me dates for Rodda's during the weeks of May 14 and 21st?

Thanks for your cooperation.

Very truly yours,

Joseph H. Zwicker

cc: Gregory D. Phillips, Esq. (by electronic and regular mail)

Peter E. Gelhaar, Esq. (by regular mail)

Michael S. D'Orsi, Esq. (by regular mail)

Brian A. Davis, Esq.

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